Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
Ľ1	24208	vascula\$10 near3 (permeab\$8 or leak\$6) or edema\$	US-PGPUB; USPAT	OR	OFF	2005/08/09 10:12
L2	12413	SrC	US-PGPUB; USPAT	OR	OFF	2005/08/09 10:13
(B)	38	1 same 2	US-PGPUB; USPAT	OR	OFF	2005/08/09 10:13
L4	1200	2 near10 (inhibit\$8 or decreas\$8)	US-PGPUB; USPAT	OR	OFF	2005/08/09 10:25
(I)	164	1 and 4	US-PGPUB; USPAT	OR	OFF	2005/08/09 10:25

3/29/00

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* * * * * * * STN Columbus * * *
FILE 'HOME' ENTERED AT 11:16:39 ON 09 AUG 2005
=> fil .bec
COST IN U.S. DOLLARS
                                                   SINCE FILE
                                                                    TOTAL
                                                        ENTRY
                                                                  SESSION
                                                         0.21
                                                                     0.21
FULL ESTIMATED COST
FILES 'MEDLINE, SCISEARCH, LIFESCI, BIOTECHDS, BIOSIS, EMBASE, HCAPLUS, NTIS,
       ESBIOBASE, BIOTECHNO, WPIDS' ENTERED AT 11:16:51 ON 09 AUG 2005
ALL COPYRIGHTS AND RESTRICTIONS APPLY. SEE HELP USAGETERMS FOR DETAILS.
11 FILES IN THE FILE LIST
=> s vascula?(3a)(permeab? or leak?) or edema?
FILE 'MEDLINE'
        377870 VASCULA?
         99161 PERMEAB?
         40828 LEAK?
          7534 VASCULA? (3A) (PERMEAB? OR LEAK?)
         81762 EDEMA?
         87886 VASCULA? (3A) (PERMEAB? OR LEAK?) OR EDEMA?
L1
FILE 'SCISEARCH'
        238013 VASCULA?
        107218 PERMEAB?
         53800 LEAK?
          8557 VASCULA? (3A) (PERMEAB? OR LEAK?)
         38623 EDEMA?
         45885 VASCULA? (3A) (PERMEAB? OR LEAK?) OR EDEMA?
L2
FILE 'LIFESCI'
         25385 VASCULA?
         18131 PERMEAB?
          5687 LEAK?
          1231 VASCULA? (3A) (PERMEAB? OR LEAK?)
          4426 EDEMA?
          5488 VASCULA? (3A) (PERMEAB? OR LEAK?) OR EDEMA?
L3
FILE 'BIOTECHDS'
          4388 VASCULA?
          3346 PERMEAB?
          1010 LEAK?
           142 VASCULA? (3A) (PERMEAB? OR LEAK?)
           372 EDEMA?
L4
           501 VASCULA? (3A) (PERMEAB? OR LEAK?) OR EDEMA?
FILE 'BIOSIS'
       2176595 VASCULA?
         95570 PERMEAB?
         35473 LEAK?
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9299 VASCULA? (3A) (PERMEAB? OR LEAK?)

6974 VASCULA? (3A) (PERMEAB? OR LEAK?)

66709 VASCULA? (3A) (PERMEAB? OR LEAK?) OR EDEMA?

91599 VASCULA? (3A) (PERMEAB? OR LEAK?) OR EDEMA?

59036 EDEMA?

379562 VASCULA? 84610 PERMEAB? 37526 LEAK?

86100 EDEMA?

L5

L6

FILE 'EMBASE'

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FILE 'HCAPLUS'
       153048 VASCULA?
        212758 PERMEAB?
         88247 LEAK?
          6802 VASCULA? (3A) (PERMEAB? OR LEAK?)
         31053 EDEMA?
L7
         36638 VASCULA? (3A) (PERMEAB? OR LEAK?) OR EDEMA?
FILE 'NTIS'
         2551 VASCULA?
         12986 PERMEAB?
         15104 LEAK?
            94 VASCULA? (3A) (PERMEAB? OR LEAK?)
           803 EDEMA?
L8
           880 VASCULA? (3A) (PERMEAB? OR LEAK?) OR EDEMA?
FILE 'ESBIOBASE'
         75800 VASCULA?
         54330 PERMEAB?
         10529 LEAK?
          2372 VASCULA? (3A) (PERMEAB? OR LEAK?)
          7737 EDEMA?
          9776 VASCULA? (3A) (PERMEAB? OR LEAK?) OR EDEMA?
L9
FILE 'BIOTECHNO'
         33459 VASCULA?
         18106 PERMEAB?
          4574 LEAK?
          1268 VASCULA? (3A) (PERMEAB? OR LEAK?)
          3864 EDEMA?
          4950 VASCULA? (3A) (PERMEAB? OR LEAK?) OR EDEMA?
L10
FILE 'WPIDS'
        23624 VASCULA?
     113129 PERMEAB?
        149364 LEAK?
           501 VASCULA? (3A) (PERMEAB? OR LEAK?)
          2903 EDEMA?
          3332 VASCULA? (3A) (PERMEAB? OR LEAK?) OR EDEMA?
L11
TOTAL FOR ALL FILES
      353644 VASCULA? (3A) (PERMEAB? OR LEAK?) OR EDEMA?
L12
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FILE 'MEDLINE'
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FILE 'SCISEARCH'
L14
      15142 SRC
FILE 'LIFESCI'
L15
      5826 SRC
FILE 'BIOTECHDS'
          318 SRC
FILE 'BIOSIS'
       15137 SRC
FILE 'EMBASE'
        11671 SRC
FILE 'HCAPLUS'
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L19

15623 SRC

FILE 'NTIS'

L20 2030 SRC

FILE 'ESBIOBASE'

L21 8142 SRC

FILE 'BIOTECHNO'

L22 7046 SRC

FILE 'WPIDS'

L23 976 SRC

TOTAL FOR ALL FILES

L24 98131 SRC

=> s 112 and 124

FILE 'MEDLINE'

L25 43 L1 AND L13

FILE 'SCISEARCH'

L26 54 L2 AND L14

FILE 'LIFESCI'

L27 11 L3 AND L15

FILE 'BIOTECHDS'

L28 3 L4 AND L16

FILE 'BIOSIS'

L29 50 L5 AND L17

FILE 'EMBASE'

L30 34 L6 AND L18

FILE 'HCAPLUS'

L31 71 L7 AND L19

FILE 'NTIS'

L32 1 L8 AND L20

FILE 'ESBIOBASE'

L33 26 L9 AND L21

FILE 'BIOTECHNO'

L34 9 L10 AND L22

FILE 'WPIDS'

L35 39 L11 AND L23

TOTAL FOR ALL FILES

L36 341 L12 AND L24

=> dup rem 136

PROCESSING COMPLETED FOR L36

L37 153 DUP REM L36 (188 DUPLICATES REMOVED)

=> d 1-30

L37 ANSWER 1 OF 153 BIOTECHDS COPYRIGHT 2005 THE THOMSON CORP. on STN

TI Use of pasireotide in manufacturing a medicament for treating disorders of growth regulation, acromegaly, diabetes, obesity, Grave's disease, macular edema, cancer and sleep apnea in a selected patient population;

pasirotide and gene expression profiling for use in disease therapy

- AU SAULNIER M
- AN 2005-19896 BIOTECHDS
- PI WO 2005053732 16 Jun 2005
- L37 ANSWER 2 OF 153 HCAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 2
- TI Preparation of imidazol[1,5-a]pyrazine derivatives, pharmaceutical compositions containing them, and their uses for prevention or treatment of protein tyrosine kinase-related diseases
- SO Jpn. Kokai Tokkyo Koho, 75 pp.

CODEN: JKXXAF

- IN Mukoyama, Harunobu; Nishimura, Toshihiro; Nakayama, Akiko; Kikuchi, Shinji; Komatsu, Yoshimitsu; Onoda, Hideki
- AN 2005:299462 HCAPLUS
- DN 142:355287

ΡI

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2005089352	A2	20050407	JP 2003-323350	20030916

- L37 ANSWER 3 OF 153 HCAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 3
- TI Pyrazolo[1,5-a]pyrimidine derivatives, prophylactic or therapeutic agents containing them for protein tyrosine kinase-related diseases, and combination drugs containing them
- SO Jpn. Kokai Tokkyo Koho, 80 pp.
- CODEN: JKXXAF
 IN Mukovama. Harunobu: Shiohara. Hiroaki: Nishim
- IN Mukoyama, Harunobu; Shiohara, Hiroaki; Nishimura, Toshihiro; Nakayama, Akiko; Kikuchi, Shinji; Komatsu, Yoshimitsu; Onoda, Hideki
- AN 2005:33492 HCAPLUS
- DN 142:127563

	PATENT NO.	KIND DATE		APPLICATION NO.	DATE	
						
ΡI	JP 2005008581	A2	20050113	JP 2003-175930	20030620	

- L37 ANSWER 4 OF 153 WPIDS COPYRIGHT 2005 THE THOMSON CORP on STN
- TI New thieno/furo-pyridine derivatives are protein tyrosine kinase inhibitors useful for treatment of e.g. ocular disorders, cardiovascular disorders, cancer.
- PI WO 2005010009 Al 20050203 (200515)* EN 391 C07D495-04
 - RW: AT BE BG BW CH CY CZ DE DK EA EE ES FI FR GB GH GM GR HU IE IT KE LS LU MC MW MZ NA NL OA PL PT RO SD SE SI SK SL SZ TR TZ UG ZM ZW
 - W: AE AG AL AM AT AU AZ BA BB BG BR BW BY BZ CA CH CN CO CR CU CZ DE DK DM DZ EC EE EG ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NA NI NO NZ OM PG PH PL PT RO RU SC SD SE SG SK SL SY TJ TM TN TR TT TZ UA UG US UZ VC VN YU ZA ZM ZW
 - US 2005026944 A1 20050203 (200515) C07D491-02
- IN BETSCHMANN, P; BURCHAT, A F; CALDERWOOD, D J; CURTIN, M L; DAVIDSEN, S K; DAVIS, H M; FREY, R R; HEYMAN, H R; HIRST, G C; HRNCIAR, P; MICHAELIDES, M R; MUCKEY, M A; RAFFERTY, P; WADA, C K
- L37 ANSWER 5 OF 153 WPIDS COPYRIGHT 2005 THE THOMSON CORP on STN
- New pyrrole pyridine derivatives are protein tyrosine kinase inhibitors useful to treat e.g. cancers, ocular conditions, cardiovascular conditions, chronic inflammation, systemic lupus, psoriasis, Paget's disease, stroke and endometriosis.
- PI US 2005043347 A1 20050224 (200520) * 181 C07D491-02
- IN BETSCHMANN, P; BURCHAT, A F; CALDERWOOD, D J; CURTIN, M L; DAVIDSEN, S K; DAVIS, H M; FREY, R R; HEYMAN, H R; HIRST, G C; HRNCIAR, P; MICHAELIDES, M R; MUCKEY, M A; RAFFERTY, P; WADA, C K
- L37 ANSWER 6 OF 153 WPIDS COPYRIGHT 2005 THE THOMSON CORP on STN
- TI New thienopyridine derivatives are protein kinase inhibitors, useful for treating e.g. cancer, ocular condition, cardiovascular condition, diabetic condition, sickle cell anemia and inflammatory bowel disease.

- PI US 2005020619 A1 20050127 (200513) * 76 C07D498-02
- IN BETSCHMANN, P; BURCHAT, A; CALDERWOOD, D; CURTIN, M L; DAVIDSEN, S K; DAVIS, H M; FREY, R R; HEYMAN, H R; HIRST, G; HRNCIAR, P; MICHAELIDES, M; RAFFERTY, P
- L37 ANSWER 7 OF 153 SCISEARCH COPYRIGHT (c) 2005 The Thomson Corporation on STN
- TI HIF-1 alpha, STAT3, CBP/p300 and Ref-1/APE are components of a transcriptional complexthat regulates **Src**-dependent hypoxia-induced expression of VEGF in pancreatic and prostate carcinomas
- SO ONCOGENE, (28 APR 2005) Vol. 24, No. 19, pp. 3110-3120. ISSN: 0950-9232.
- AU Gray M J; Zhang J; Ellis L M; Semenza G L; Evans D B; Watowich S S; Gallick G E (Reprint)
- AN 2005:472725 SCISEARCH
- L37 ANSWER 8 OF 153 MEDLINE on STN DUPLICATE 4
- TI Tumor metastasis but not tumor growth is dependent on **Src** -mediated **vascular permeability**.
- SO Blood, (2005 Feb 15) 105 (4) 1508-14. Electronic Publication: 2004-10-14. Journal code: 7603509. ISSN: 0006-4971.
- AU Criscuoli Michele L; Nguyen Mai; Eliceiri Brian P
- AN 2005060850 MEDLINE
- L37 ANSWER 9 OF 153 SCISEARCH COPYRIGHT (c) 2005 The Thomson Corporation on STN
- TI Conditional knockout of focal adhesion kinase in endothelial cells reveals its role in angiogenesis and vascular development in late embryogenesis
- SO JOURNAL OF CELL BIOLOGY, (20 JUN 2005) Vol. 169, No. 6, pp. 941-952. ISSN: 0021-9525.
- AU Shen T L; Park A Y J; Alcaraz A; Peng X; Jang I; Koni P; Flavell R A; Gu H; Guan J L (Reprint)
- AN 2005:663724 SCISEARCH
- L37 ANSWER 10 OF 153 MEDLINE on STN DUPLICATE 5
- TI Inhibition of the **Src** and Jak Kinases Protects against Lipopolysaccharide-induced Acute Lung Injury.
- SO American journal of respiratory and critical care medicine, (2005 Apr 15) 171 (8) 858-67. Electronic Publication: 2005-01-21.

 Journal code: 9421642. ISSN: 1073-449X.
- AU Severgnini Mariano; Takahashi Satoe; Tu Powen; Perides George; Homer Robert J; Jhung Jhung W; Bhavsar Deepa; Cochran Brent H; Simon Amy R
- AN 2005184113 IN-PROCESS
- L37 ANSWER 11 OF 153 MEDLINE ON STN DUPLICATE 6
- TI Vascular endothelial-cadherin tyrosine phosphorylation in angiogenic and quiescent adult tissues.
- SO Circulation research, (2005 Feb 18) 96 (3) 384-91. Electronic Publication: 2005-01-20.

 Journal code: 0047103. ISSN: 1524-4571.
- AU Lambeng Nathalie; Wallez Yann; Rampon Christine; Cand Francine; Christe Georges; Gulino-Debrac Danielle; Vilgrain Isabelle; Huber Philippe
- AN 2005088277 MEDLINE
- L37 ANSWER 12 OF 153 EMBASE COPYRIGHT 2005 ELSEVIER INC. ALL RIGHTS RESERVED. ON STN
- TI Molecularly targeted therapy for gastrointestinal cancer.
- SO Current Cancer Drug Targets, (2005) Vol. 5, No. 3, pp. 171-193.
 Refs: 102
 ISSN: 1568-0096 CODEN: CCDTB
- AU Wiedmann M.W.; Caca K.
- AN 2005226075 EMBASE

- TI Modulation of peripheral inflammation in sensory ganglia by nuclear factor (kappa)B decoy oligodeoxynucleotide: involvement of **SRC** kinase pathway.
- SO Neuroscience letters, (2005 Jun 10-17) 381 (1-2) 114-9. Electronic Publication: 2005-02-25.

 Journal code: 7600130. ISSN: 0304-3940.
- AU Iqwe Orisa J
- AN 2005245223 IN-PROCESS
- L37 ANSWER 14 OF 153 SCISEARCH COPYRIGHT (c) 2005 The Thomson Corporation on STN
- TI Comparison of the signaling mechanisms by which VEGF, $\dot{H}2O2$, and phosphatase inhibitors activate endothelial cell ERK1/2 MAP-kinase
- SO MICROVASCULAR RESEARCH, (JAN 2005) Vol. 69, No. 1-2, pp. 36-44. ISSN: 0026-2862.
- AU Tao Q; Spring S C; Terman B I (Reprint)
- AN 2005:428794 SCISEARCH
- ·L37 ANSWER 15 OF 153 SCISEARCH COPYRIGHT (c) 2005 The Thomson Corporation on STN
- TI 17 beta-estradiol transiently disrupts adherens junctions in endothelial cells
- SO FASEB JOURNAL, (MAY 2005) Vol. 19, No. 7. ISSN: 0892-6638.
- AU Groten T (Reprint); Pierce A A; Huen A C; Schnaper H W
- AN 2005:613823 SCISEARCH
- L37 ANSWER 16 OF 153 BIOTECHDS COPYRIGHT 2005 THE THOMSON CORP. on STN

 TI Composition for treating or preventing a blood-brain barrier disorder,
 e.g. multiple sclerosis, ischemia, Alzheimer's disease, brain tumors, or
 hypertension, comprises an src-suppressed C kinase substrate
 protein:
 - involving vector-mediated gene transfer and expression in host cell for therapy
- AU KIM K; LEE S; KIM W
- AN 2005-03352 BIOTECHDS
- PI WO 2004110476 23 Dec 2004
- L37 ANSWER 17 OF 153 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN DUPLICATE 9
- TI Methods and compositions useful for modulation of angiogenesis and vascular permeability using SRC or Yes tyrosine kinases.
- SO Official Gazette of the United States Patent and Trademark Office Patents, (Feb 3 2004) Vol. 1279, No. 1. http://www.uspto.gov/web/menu/patdata.html.e-file.
- ISSN: 0098-1133 (ISSN print).
- AU Cheresh, David A. [Inventor, Reprint Author]; Eliceiri, Brian [Inventor]
- AN 2004:130012 BIOSIS
- L37 ANSWER 18 OF 153 HCAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 10
- TI Preparation of 4-[(2,4-dichloro-5-methoxyphenyl)amino]-6-alkoxy-3-quinolinecarbonitriles as **Src** inhibitors for the treatment of ischemic injury
- SO PCT Int. Appl., 43 pp. CODEN: PIXXD2
- IN Boschelli, Diane Harris; Zaleska, Margaret Maria; Boschelli, Frank Charles; Arndt, Kim Timothy
- AN 2004:740166 HCAPLUS
- DN 141:243354
 - PATENT NO. KIND DATE APPLICATION NO. DATE
- PI WO 2004075898 A1 20040910 WO 2004-US4904 20040219 W: AE, AE, AG, AL, AL, AM, AM, AM, AT, AT, AU, AZ, AZ, BA, BB, BG,

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              MZ, MZ, NA, NI
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                                   20041118
                                                 US 2004-780973
                                                                          20040218
     US 2004229880
                            A1
     ANSWER 19 OF 153 HCAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 11
L37
TI
     Inhibition of src family kinases for the treatment of
     reperfusion injury related to revascularization
SO
     PCT Int. Appl., 62 pp.
     CODEN: PIXXD2
IN
     Losordo, Douglas W.
     2004:331894 HCAPLUS
AN
DN
     140:350577
                                                                          DATE
                                   DATE
                                                APPLICATION NO.
     PATENT NO.
                           KIND
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                                                WO 2003-US31430
     WO 2004032709
                            Α2
                                    20040422
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PI
     WO 2004032709
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                                   20041007
              AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
              CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
              GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
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                                    20040422
                                                CA 2003-2500368
                                                                          20031003
      CA 2500368
                             AA
     ANSWER 20 OF 153 HCAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 12
L37
     Preparation of vasculostatic agents and methods of use
TI
     PCT Int. Appl., 230 pp.
SO
     CODEN: PIXXD2
     Wrasidlo, Wolfgang; Doukas, John; Royston, Ivor; Noronha, Glenn; Hood,
ΙN
     John D.; Dneprovskaia, Elena; Gong, Xianchang; Splittgerber, Ute; Zhao,
     Ningning
AN
     2004:308364 HCAPLUS
     140:321386
DN
      PATENT NO.
                            KIND
                                    DATE
                                                 APPLICATION NO.
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     WO 2004030635
                            A2
                                    20040415
                                                 WO 2003-US31721
                                                                          20031002
PΙ
     WO 2004030635
                            A3
                                    20040812
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              CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE,
              GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK,
              LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ,
              OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM,
              TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
          RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
              KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES,
              FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR,
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     CA 2500727
                                    20040415
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                                                                          20031002
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     US 2004167198
                                    20040826
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                             A1
                                    20050706
                                                 EP 2003-774610
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     EP 1549614
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          R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
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BG, BR, BR, BW, BY, BY, BZ, BZ, CA, CH, CN, CN, CO, CO, CR, CR,

- L37 ANSWER 21 OF 153 WPIDS COPYRIGHT 2005 THE THOMSON CORP on STN
- TI New pyrrolotriazine derivatives useful for treatment of proliferative disease e.g. cancer, inflammation and autoimmune disease.
- PI WO 2004013145 A1 20040212 (200420)* EN 71 C07D487-04
 - RW: AT BE BG CH CY CZ DE DK EA EE ES FI FR GB GH GM GR HU IE IT KE LS LU MC MW MZ NL OA PT RO SD SE SI SK SL SZ TR TZ UG ZM ZW
 - W: AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ DE DK DM DZ EC EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NI NO NZ OM PG PH PL PT RO RU SC SD SE SG SK SL SY TJ TM TN TR TT TZ UA UG US UZ VC VN YU ZA ZM ZW
 - US 2004063708 A1 20040401 (200425) A61K031-53 AU 2003265349 A1 20040223 (200453) C07D487-04 EP 1543009 A1 20050622 (200541) EN C07D487-04
 - R: AL AT BE BG CH CY CZ DE DK EE ES FI FR GB GR HU IE IT LI LT LU LV MC MK NL PT RO SE SI SK TR
- IN BHIDE, R S; BORZILLERI, R M
- L37 ANSWER 22 OF 153 LIFESCI COPYRIGHT 2005 CSA on STN
- TI Activation of Vascular Endothelial Growth Factor Receptor-3 and Its Downstream Signaling Promote Cell Survival under Oxidative Stress
- SO Journal of Biological Chemistry [J. Biol. Chem.], (20040625) vol. 279, no. 26, pp. 27088-27097.

 ISSN: 0021-9258.
- AU Wang, J.F.; Zhang, X.; Groopman, J.E.
- AN 2004:85045 LIFESCI
- L37 ANSWER 23 OF 153 SCISEARCH COPYRIGHT (c) 2005 The Thomson Corporation on STN
- TI Angiogenesis and its role in the behavior of astrocytic brain tumors
- SO FRONTIERS IN BIOSCIENCE, (1 SEP 2004) Vol. 9, Supp. [S], pp. 3105-3123. ISSN: 1093-9946.
- AU Stiver S I (Reprint)
- AN 2004:819609 SCISEARCH
- L37 ANSWER 24 OF 153 MEDLINE on STN DUPLICATE 13
- TI **Src** family kinase inhibitor PP1 reduces secondary damage after spinal cord compression in rats.
- SO Journal of neurotrauma, (2004 Jul) 21 (7) 923-31. Journal code: 8811626. ISSN: 0897-7151.
- AU Akiyama Chihiro; Yuguchi Takamichi; Nishio Masami; Tomishima Takahiro; Fujinaka Toshiyuki; Taniguchi Masaaki; Nakajima Yoshikazu; Kohmura Eiji; Yoshimine Toshiki
- AN 2004404797 MEDLINE
- L37 ANSWER 25 OF 153 MEDLINE ON STN DUPLICATE 14
- TI Signaling pathways for early brain injury after subarachnoid hemorrhage.
- SO Journal of cerebral blood flow and metabolism : official journal of the International Society of Cerebral Blood Flow and Metabolism, (2004 Aug) 24 (8) 916-25.
 - Journal code: 8112566. ISSN: 0271-678X.
- AU Kusaka Gen; Ishikawa Mami; Nanda Anil; Granger D Neil; Zhang John H
- AN 2004454757 MEDLINE
- L37 ANSWER 26 OF 153 MEDLINE ON STN DUPLICATE 15
- TI Src blockade stabilizes a Flk/cadherin complex, reducing edema and tissue injury following myocardial infarction.
- SO Journal of clinical investigation, (2004 Mar) 113 (6) 885-94. Journal code: 7802877. ISSN: 0021-9738.
- AU Weis Sara; Shintani Satoshi; Weber Alberto; Kirchmair Rudolf; Wood Malcolm; Cravens Adrianna; McSharry Heather; Iwakura Atsushi; Yoon Young-Sup; Himes Nathan; Burstein Deborah; Doukas John; Soll Richard;

Losordo Douglas; Cheresh David

- AN 2004172915 MEDLINE
- L37 ANSWER 27 OF 153 MEDLINE on STN DUPLICATE 16
- TI Angiogenic signal triggered by ischemic stress induces myocardial repair in rat during chronic infarction.
- SO Journal of molecular and cellular cardiology, (2004 Apr) 36 (4) 547-59. Journal code: 0262322. ISSN: 0022-2828.
- AU Fukuda Shoji; Kaga Shigeaki; Sasaki Hiroaki; Zhan Lijun; Zhu Li; Otani Hajime; Kalfin Reni; Das Dipak K; Maulik Nilanjana
- AN 2004186500 MEDLINE
- L37 ANSWER 28 OF 153 SCISEARCH COPYRIGHT (c) 2005 The Thomson Corporation on STN
- TI Matrix-specific activation of **Src** and Rho initiates capillary morphogenesis of endothelial cells
- SO FASEB JOURNAL, (MAR 2004) Vol. 18, No. 3, pp. 457-468. ISSN: 0892-6638.
- AU Liu Y Q; Senger D R (Reprint)
- AN 2004:311780 SCISEARCH
- L37 ANSWER 29 OF 153 MEDLINE on STN DUPLICATE 17
- TI Influenza virus inhibits ENaC and lung fluid clearance.
- SO American journal of physiology. Lung cellular and molecular physiology, (2004 Aug) 287 (2) L366-73. Electronic Publication: 2004-04-30. Journal code: 100901229. ISSN: 1040-0605.
- AU Chen Xi-Juan; Seth Shaguna; Yue Gang; Kamat Pradip; Compans Richard W; Guidot David; Brown Lou Ann; Eaton Douglas C; Jain Lucky
- AN 2004342287 MEDLINE
- L37 ANSWER 30 OF 153 Elsevier BIOBASE COPYRIGHT 2005 Elsevier Science B.V. on STN
- AN 2004184814 ESBIOBASE
- TI Influenza virus inhibits ENaC and lung fluid clearance
- AU Chen X.-J.; Seth S.; Yue G.; Kamat P.; Compans R.W.; Guidot D.; Brown L.A.; Eaton D.C.; Jain L.
- CS L. Jain, Dept. of Pediatrics, Emory Univ. School of Medicine, 2040 Ridgewood Dr., NE, Atlanta, GA 30322, United States. E-mail: Ljain@emory.edu
- SO American Journal of Physiology Lung Cellular and Molecular Physiology, (2004), 287/2 31-2 (L366-L373), 34 reference(s)

 CODEN: APLPE7 ISSN: 1040-0605
- DT Journal; Article
- CY United States
- LA English
- SL English,

=> d ab 5,8,24,26

- L37 ANSWER 5 OF 153 WPIDS COPYRIGHT 2005 THE THOMSON CORP on STN
- AB US2005043347 A UPAB: 20050324

NOVELTY - Pyrrole pyridine derivatives (I) and their salts are new.

DETAILED DESCRIPTION - Pyrrole pyridine derivatives of formula (I) and their salts are new.

X = 0 or S;

Z = C or N;

R1 = H, alkenyl, alkoxyalkynyl, alkoxycarbonyl, alkoxycarbonylalkenyl, alkoxycarbonylalkyl, alkoxycarbonylalkynyl, alkyl, alkynyl, arylalkenyl, arylalkyl, arylalkynyl, aryloxyalkynyl, arylsulfanylalkyl, arylsulfanylalkynyl, arylsulfonyloxyalkenyl, carboxy, carboxyalkenyl, carboxyalkynyl, carboxyalkynyl, cyanoalkyl, cyanoalkyl, cyanoalkynyl, cycloalkylalkoxyalkynyl, cycloalkylalkenyl, cycloalkylalkynyl,

formylalkenyl, formylalkyl, halo, haloalkyl, heteroaryl, heteroarylalkenyl, heteroarylalkyl, heteroarylalkynyl, heteroarylcarbonyl, heteroarylcarbonylalkenyl, heteroarylcarbonylalkyl, heterocyclyl, heterocyclylalkenyl, heterocyclylalkyl, heterocyclylalkylcarbonyl, heterocyclylalkynyl, heterocyclylcarbonyl, heterocyclylcarbonylalkenyl, heterocyclylcarbonylalkenyl, hydroxyalkenyl, hydroxyalkenyl, hydroxyalkynyl, NR-aR-b, (NR-aR-b)carbonyl, (NR-aR-b)(carbonyl)alkenyl, (NR-aR-b)(carbonyl)alkynyl, (nitro)alkenyl, nitro alkyl or nitroalkynyl;

R2 = H, alkyl or absent;

R3 = (hetero)aryl, heterocyclyl (both optionally substituted with 1-3 of alkoxy, alkyl, CN, halo, haloalkoxy, haloalkyl, (hetero)aryl, heterocyclyl, OH, hydroxyalkyl, LR4 or NR-aR-b) or halo;

 $L = O, (CH2)mC(0)NR5, NR5C(0)(CH2)m, NR5SO2, SO2NR5, \\ (CH2)mN(R5)C(0)N(R6)(CH2)n or (CH2)mN(R5)C(S)N(R6)(CH2)n (where each group is drawn with its right end attached to R4);$

R4 = cycloalkyl, (hetero)aryl, (hetero)arylalkyl or heterocyclyl(alkyl);

R5, R6 = H or alkyl;

R-a, R-b = arylalkyl, aryl (both substituted with 1-5 of alkoxy, alkyl, CN, halo, haloalkoxy, haloalkyl, nitro or oxo), arylalkoxycarbonyl, arylalkoxycarbonylalkyl, arylcarbonyl, arylsulfonyl, cycloalkyl, cycloalkyl, heteroaryl, heteroarylalkyl, heteroarylcarbonyl, heterocyclyl, heterocyclylalkyl, heterocyclylcarbonyl (all optionally substituted with 1-5 of alkenyl, alkoxy, alkoxycarbonyl, alkyl, alkylcarbonyl, aryl, arylalkyl, halo, haloalkoxy, haloalkyl, OH, nitro, NR-cR-d, (NR-cR-d)alkyl, (NR-cR-d)alkylcarbonyl, (NR-cR-d)carbonyl, (NR-cR-d)carbonylalkyl, oxo or spiroheterocyclyl), H, alkenyl, alkoxyalkyl, alkoxycarbonyl, alkoxycarbonylalkyl, alkyl, alkylcarbonyl, alkylsulfanylalkyl, alkylsulfonyl, carboxyalkyl, formylalkyl, heteroarylsulfonyl, heterocyclylalkylcarbonyl, heterocyclylsulfonyl, hydroxyalkyl, hydroxyalkyl, (NR-cR-d)alkyl, (NR-cR-d)alkylcarbonyl, (NR-cR-d)carbonyl or (NR-cR-d)carbonylalkyl;

R-c, R-d = (hetero)aryl, heterocyclyl, (both optionally substituted with 1-5 of alkenyl, alkoxy, alkyl, halo, haloalkoxy, haloalkyl, OH or NO2), H, alkoxy, alkyl, carboxyalkyl, cycloalkyl, haloalkyl, heterocyclylalkyl, hydroxyalkoxyalkyl, hydroxyalkyl or (NR-eR-f)alkyl;

R-e, R-f = H or alkyl; and m, n = 0-1.

Provided that at least two of the three substituents are other than ${\tt LR4}\,.$

ACTIVITY - Cytostatic; Ophthalmological; Cardiovascular-Gen.; Antidiabetic; Antianemic; Antisickling; Antiinflammatory; Dermatological; Immunosuppressive; Nephrotropic; Gastrointestinal-Gen.; Antiarthritic; Antirheumatic; Osteopathic; Neuroprotective; Antibacterial; Antipsoriatic; Gynecological; Hepatotropic; Antithyroid; Respiratory-Gen.; Antiasthmatic; Vulnerary; Antiemetic; Cerebroprotective; Vasotropic; Virucide; Anti-HIV; Protozoacide; Muscular-Gen.; Anticonvulsant; Hypotensive.

MECHANISM OF ACTION - Protein tyrosine kinase inhibitor. (I) were tested for their kinase insert domain containing receptor (KDR) tyrosine kinase inhibitory activity using complimentary DNAs isolated from HUVEC cells. The results showed that the median inhibitory concentration of (I) was 0.002-50 micro M.

USE - (I) are useful to inhibit one or more protein kinases (KDR, lymphocyte specific protein tyrosine kinase (Lck) (both preferred), Ckit, colony stimulating factor (CSF)-1R, platelet derivative growth factor receptor (PDGFR) beta, PDGFR alpha, Flt-1, Flt-3, Flt-4, Tie-2, Src, Fyn, Lyn, Blk, Hck, Fgr, Cot and Yes) and to treat cancers (preferred), ocular conditions, cardiovascular conditions, Crow-Fukase (POEMS) syndrome, diabetic conditions, sickle cell anemia, chronic inflammation, systemic lupus, glomerulonephritis, synovitis, inflammatory bowel disease, Crohn's disease, rheumatoid arthritis, osteoarthritis, multiple sclerosis, graft rejection, lyme disease, sepsis, von Hippel Lindau disease, pemphigoid, psoriasis, Paget's disease, polycystic kidney

disease, fibrosis, sarcoidosis, cirrhosis, thyroiditis, hyperviscosity syndrome, Osler-Weber-Rendu disease, chronic occlusive pulmonary disease, asthma or edema following bums, trauma, radiation, stroke, hypoxia, ischemia, ovarian hyperstimulation syndrome, preeclampsia, menometrorrhagia, endometriosis or infection by Herpes simplex, Herpes Zoster, human immunodeficiency virus diseases, parapoxyvirus, protozoa infection and toxoplasmosis (claimed). (I) are also useful to treat benign and neoplastic proliferative diseases (human cancers of e.g. lung and breast), disorders of the immune system such as autoimmune diseases (e.g. thyroiditis, type 1 diabetes, multiple sclerosis, sarcoidosis, inflammatory bowel disease, Crohn's disease, myasthenia gravis and systemic lupus erythematosus), psoriasis, organ transplant rejection (e.g. kidney rejection, graft versus host disease), hematopoietic malignancies (leukemia and lymphoma), glioblastoma, infantile hemangioma, diseases involving inappropriate vascularization (e.g. diabetic retinopathy, retinopathy of prematurity, choroidal neovascularization due to age-related macular degeneration and infantile hemangiomas), vascular endothelial growth factor mediated edema, ascites, effusions, exudates (e.g. macular edema, cerebral edema, acute lung injury and adult respiratory distress syndrome) and pulmonary hypertension (particularly thromboembolic disease).

ADVANTAGE - (I) has no toxicity, irritation and allergic responses. Dwg.0/0

- MEDLINE on STN DUPLICATE 4 1.37 ANSWER 8 OF 153 Vascular endothelial growth factor (VEGF) - induced vascular permeability (VP) is a hallmark of tumor growth and metastasis. Previous studies have shown a requirement for Src kinase in VEGF-mediated VP and signaling in blood vessels. In this study, we have examined the effect of Src-mediated reduced VP on tumor growth and metastasis. The growth and spontaneous metastasis of VEGF-expressing tumor cells were determined in Src-knockout (src(-/-)) or control mice (src(+/+)) or src(+/-). In comparison to control mice, src-null mice had a significant reduction in tumor-induced VP as well as a subsequent reduction in spontaneous metastasis. In contrast, primary tumor weight and vascular density were unchanged between src-null and control mice. Consistent with a role for Src in the extravasation of tumor cells from the circulation, direct intravenous injection of lung carcinoma cells resulted in a more than 2-fold reduction in lung tumor burden in src-null mice compared to control mice. The comparison of the results from the experimental metastasis and the spontaneous metastasis models suggests that there are defects in VP in the primary site of Src -deficient mice and that there may be an essential role for Src and Src-mediated VP in tumor metastasis to the lung.
- MEDLINE on STN **DUPLICATE 13** L37 ANSWER 24 OF 153 The synthetic pyrazolopyrimidine, 4-amino-5-(4-methylphenyl)-7-(t-AB butyl)pyrazolo[3,4-d]pyrimidine (PP1) is a novel, potent, and selective inhibitor of Src family tyrosine kinases. Vascular permeability appears to be mediated by vascular endothelial growth factor (VEGF), which requires the activation of downstream Src family kinases to exert its function. This study investigates the effects of PP1 on vascular permeability and inflammatory response in a rat spinal cord compression model. Ten minutes after compression, PP1 (PP1 group) or the vehicle only (control group) was administered. On days 1, 3, and 7 after compression, the spinal cords were removed and examined histopathologically to determine the expression of VEGF and the extent of edema and inflammation. The dryweight method was used to measure the water content of the spinal cords. mRNA levels of tumor necrosis factor a (TNFalpha) and interleukin 1beta (IL-1beta), which is related to inflammatory responses, were measured with a real-time polymerase chain reaction (RT-PCR) system 6 h after compression. Although VEGF expression was similar in both groups, the

extent of contusional lesion in the PP1 group was reduced by approximately 35% on day 3. Moreover, the water content on days 1, 3, and 7 was significantly reduced and macrophage infiltration on days 3 and 7 was dramatically reduced in the PP1 group. TNF and IL-1beta mRNA expression in the PP1 group were also significantly reduced. These results indicate that PP1 reduces secondary damage after spinal cord injury.

L37 ANSWER 26 OF 153 MEDLINE on STN Ischemia resulting from myocardial infarction (MI) promotes VEGF AΒ expression, leading to vascular permeability (VP) and edema, a process that we show here contributes to tissue injury throughout the ventricle. This permeability/edema can be assessed noninvasively by MRI and can be observed at the ultrastructural level as gaps between adjacent endothelial cells. Many of these gaps contain activated platelets adhering to exposed basement membrane, reducing vessel patency. Following MI, genetic or pharmacological blockade of Src preserves endothelial cell barrier function, suppressing VP and infarct volume, providing long-term improvement in cardiac function, fibrosis, and survival. To our surprise, an intravascular injection of VEGF into healthy animals, but not those deficient in Src, induced similar endothelial gaps, VP, platelet plugs, and some myocyte damage. Mechanistically, we show that quiescent blood vessels contain a complex involving Flk, VE-cadherin, and beta-catenin that is transiently disrupted by VEGF injection. Blockade of Src prevents disassociation of this complex with the same kinetics with which it prevents VEGF-mediated VP/edema. These findings define a molecular mechanism to account for the Src requirement in VEGF-mediated permeability and provide a basis for Src inhibition as a therapeutic option for patients with acute MI.

=> s aminopyrazolopyrimidine? or pyrazolopyrimidine? or (aminopyrazolo or pyrazolo) (3w) pyrimidine?

FILE 'MEDLINE'

- 50 AMINOPYRAZOLOPYRIMIDINE?
- 94 PYRAZOLOPYRIMIDINE?
- 141 AMINOPYRAZOLO
- 990 PYRAZOLO
- 29241 PYRIMIDINE?
 - 575 (AMINOPYRAZOLO OR PYRAZOLO) (3W) PYRIMIDINE?
- 666 AMINOPYRAZOLOPYRIMIDINE? OR PYRAZOLOPYRIMIDINE? OR (AMINOPYRAZOL O OR PYRAZOLO) (3W) PYRIMIDINE?

FILE 'SCISEARCH'

L38

- 35 AMINOPYRAZOLOPYRIMIDINE?
- 152 PYRAZOLOPYRIMIDINE?
- 102 AMINOPYRAZOLO
- 2505 PYRAZOLO
- 21984 PYRIMIDINE?
 - 843 (AMINOPYRAZOLO OR PYRAZOLO) (3W) PYRIMIDINE?
- L39 987 AMINOPYRAZOLOPYRIMIDINE? OR PYRAZOLOPYRIMIDINE? OR (AMINOPYRAZOL O OR PYRAZOLO) (3W) PYRIMIDINE?

FILE 'LIFESCI'

- 12 AMINOPYRAZOLOPYRIMIDINE?
- 34 PYRAZOLOPYRIMIDINE?
- 14 AMINOPYRAZOLO
- 158 PYRAZOLO
- 5854 PYRIMIDINE?
 - 86 (AMINOPYRAZOLO OR PYRAZOLO) (3W) PYRIMIDINE?
- L40 121 AMINOPYRAZOLOPYRIMIDINE? OR PYRAZOLOPYRIMIDINE? OR (AMINOPYRAZOL O OR PYRAZOLO) (3W) PYRIMIDINE?

- 0 AMINOPYRAZOLOPYRIMIDINE?
- 6 PYRAZOLOPYRIMIDINE?
- 3 AMINOPYRAZOLO
- 27 PYRAZOLO
- 907 PYRIMIDINE?
- 23 (AMINOPYRAZOLO OR PYRAZOLO) (3W) PYRIMIDINE?
- L41 28 AMINOPYRAZOLOPYRIMIDINE? OR PYRAZOLOPYRIMIDINE? OR (AMINOPYRAZOL O OR PYRAZOLO) (3W) PYRIMIDINE?

FILE 'BIOSIS'

- 57 AMINOPYRAZOLOPYRIMIDINE?
- 162 PYRAZOLOPYRIMIDINE?
- 141 AMINOPYRAZOLO
- 1841 PYRAZOLO
- 21961 PYRIMIDINE?
 - 700 (AMINOPYRAZOLO OR PYRAZOLO) (3W) PYRIMIDINE?
- L42 861 AMINOPYRAZOLOPYRIMIDINE? OR PYRAZOLOPYRIMIDINE? OR (AMINOPYRAZOL O OR PYRAZOLO) (3W) PYRIMIDINE?

FILE 'EMBASE'

- 47 AMINOPYRAZOLOPYRIMIDINE?
- 216 PYRAZOLOPYRIMIDINE?
- 176 AMINOPYRAZOLO
- 2571 PYRAZOLO
- 18893 PYRIMIDINE?
 - 798 (AMINOPYRAZOLO OR PYRAZOLO) (3W) PYRIMIDINE?
- L43 984 AMINOPYRAZOLOPYRIMIDINE? OR PYRAZOLOPYRIMIDINE? OR (AMINOPYRAZOL O OR PYRAZOLO) (3W) PYRIMIDINE?

FILE 'HCAPLUS'

- 194 AMINOPYRAZOLOPYRIMIDINE?
- 1501 PYRAZOLOPYRIMIDINE?
 - 417 AMINOPYRAZOLO
- 5814 PYRAZOLO
- 63957 PYRIMIDINE?
- 1819 (AMINOPYRAZOLO OR PYRAZOLO) (3W) PYRIMIDINE?
- L44 2487 AMINOPYRAZOLOPYRIMIDINE? OR PYRAZOLOPYRIMIDINE? OR (AMINOPYRAZOL O OR PYRAZOLO) (3W) PYRIMIDINE?

FILE 'NTIS'

- 1 AMINOPYRAZOLOPYRIMIDINE?
- 1 PYRAZOLOPYRIMIDINE?
- 0 AMINOPYRAZOLO
- 5 PYRAZOLO
- 529 PYRIMIDINE?
 - 1 (AMINOPYRAZOLO OR PYRAZOLO) (3W) PYRIMIDINE?
- L45 3 AMINOPYRAZOLOPYRIMIDINE? OR PYRAZOLOPYRIMIDINE? OR (AMINOPYRAZOL O OR PYRAZOLO) (3W) PYRIMIDINE?

FILE 'ESBIOBASE'

- 2 AMINOPYRAZOLOPYRIMIDINE?
- 35 PYRAZOLOPYRIMIDINE?
- 11 AMINOPYRAZOLO
- 355 PYRAZOLO
- 5103 PYRIMIDINE?
- 160 (AMINOPYRAZOLO OR PYRAZOLO) (3W) PYRIMIDINE?
- L46 194 AMINOPYRAZOLOPYRIMIDINE? OR PYRAZOLOPYRIMIDINE? OR (AMINOPYRAZOL O OR PYRAZOLO) (3W) PYRIMIDINE?

FILE 'BIOTECHNO'

- 15 AMINOPYRAZOLOPYRIMIDINE?
- 27 PYRAZOLOPYRIMIDINE?
- 42 AMINOPYRAZOLO
- 260 PYRAZOLO

5965 PYRIMIDINE?

85 (AMINOPYRAZOLO OR PYRAZOLO) (3W) PYRIMIDINE?

L47 119 AMINOPYRAZOLOPYRIMIDINE? OR PYRAZOLOPYRIMIDINE? OR (AMINOPYRAZOL O OR PYRAZOLO) (3W) PYRIMIDINE?

FILE 'WPIDS'

2 AMINOPYRAZOLOPYRIMIDINE?

143 PYRAZOLOPYRIMIDINE?

53 AMINOPYRAZOLO

2516 PYRAZOLO

13541 PYRIMIDINE?

570 (AMINOPYRAZOLO OR PYRAZOLO) (3W) PYRIMIDINE?

L48 663 AMINOPYRAZOLOPYRIMIDINE? OR PYRAZOLOPYRIMIDINE? OR (AMINOPYRAZOL O OR PYRAZOLO) (3W) PYRIMIDINE?

TOTAL FOR ALL FILES

L49 7113 AMINOPYRAZOLOPYRIMIDINE? OR PYRAZOLOPYRIMIDINE? OR (AMINOPYRAZOL O OR PYRAZOLO)(3W) PYRIMIDINE?

=> s 149 and src(4a)inhibit?

FILE 'MEDLINE'

16220 SRC

1193234 INHIBIT?

1673 SRC(4A) INHIBIT?

L50 166 L38 AND SRC(4A) INHIBIT?

FILE 'SCISEARCH'

15142 SRC

985798 INHIBIT?

1730 SRC(4A) INHIBIT?

L51 100 L39 AND SRC(4A) INHIBIT?

FILE 'LIFESCI'

5826 SRC

323425 INHIBIT?

585 SRC(4A) INHIBIT?

L52 27 L40 AND SRC(4A)INHIBIT?

FILE 'BIOTECHDS'

318 SRC

54401 INHIBIT?

34 SRC(4A) INHIBIT?

L53 1 L41 AND SRC(4A) INHIBIT?

FILE 'BIOSIS'

15137 SRC

1269669 INHIBIT?

2000 SRC(4A) INHIBIT?

L54 98 L42 AND SRC(4A) INHIBIT?

FILE 'EMBASE'

11671 SRC

1080241 INHIBIT?

1606 SRC(4A) INHIBIT?

L55 171 L43 AND SRC(4A)INHIBIT?

FILE 'HCAPLUS'

15623 SRC

1761127 INHIBIT?

2067 SRC(4A) INHIBIT?

L56 116 L44 AND SRC(4A) INHIBIT?

FILE 'NTIS'

2030 SRC

20752 INHIBIT?

10 SRC(4A) INHIBIT?

L57

0 L45 AND SRC(4A) INHIBIT?

FILE 'ESBIOBASE'

8142 SRC

427731 INHIBIT?

1371 SRC(4A) INHIBIT?

Ŀ58

91 L46 AND SRC(4A) INHIBIT?

FILE 'BIOTECHNO'

7046 SRC

301415 INHIBIT?

754 SRC(4A) INHIBIT?

L59

38 L47 AND SRC(4A) INHIBIT?

FILE 'WPIDS'

976 SRC

236898 INHIBIT?

187 SRC(4A) INHIBIT?

L60

6 L48 AND SRC(4A) INHIBIT?

TOTAL FOR ALL FILES

L61 814 L49 AND SRC(4A) INHIBIT?

=> dup rem 161

PROCESSING COMPLETED FOR L61

1.62 280

280 DUP REM L61 (534 DUPLICATES REMOVED)

=> d 1-100

L62 ANSWER 1 OF 280 HCAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 1

TI **Pyrazolo**[1,5-a]**pyrimidine** derivatives, prophylactic or therapeutic agents containing them for protein tyrosine kinase-related diseases, and combination drugs containing them

SO Jpn. Kokai Tokkyo Koho, 80 pp.

CODEN: JKXXAF

IN Mukoyama, Harunobu; Shiohara, Hiroaki; Nishimura, Toshihiro; Nakayama, Akiko; Kikuchi, Shinji; Komatsu, Yoshimitsu; Onoda, Hideki

AN 2005:33492 HCAPLUS

DN 142:127563

PΙ

PATENT NO. KIND DATE APPLICATION NO. DATE

JP 2005008581 A2 20050113 JP 2003-175930 20030620

L62 ANSWER 2 OF 280 MEDLINE on STN

DUPLICATE 2

TI p38 kinase-mediated transactivation of the epidermal growth factor receptor is required for dedifferentiation of renal epithelial cells after oxidant injury.

SO Journal of biological chemistry, (2005 Jun 3) 280 (22) 21036-42. Electronic Publication: 2005-03-28.

Journal code: 2985121R. ISSN: 0021-9258.

AU Zhuang Shougang; Yan Yan; Han Jiahuai; Schnellmann Rick G

AN 2005279091 MEDLINE

L62 ANSWER 3 OF 280 EMBASE COPYRIGHT 2005 ELSEVIER INC. ALL RIGHTS RESERVED. on STN

TI Src-mediated tyrosine phosphorylation of p47(phox) in hyperoxia-induced activation of NADPH oxidase and generation of reactive oxygen species in lung endothelial cells.

SO Journal of Biological Chemistry, (27 May 2005) Vol. 280, No. 21, pp. 20700-20711.

Refs: 53

ISSN: 0021-9258 CODEN: JBCHA3

- AU Chowdhury A.K.; Watkins T.; Parinandi N.L.; Saatian B.; Kleinberg M.E.; Usatyuk P.V.; Natarajan V.
- AN 2005250924 EMBASE
- L62 ANSWER 4 OF 280 EMBASE COPYRIGHT 2005 ELSEVIER INC. ALL RIGHTS RESERVED. on STN
- TI Survival function of protein kinase Ct, as a novel nitrosamine 4-(methylinitrosamino) -1-(3-pyridyl)-1-butanone-activated bad kinase.
- SO Journal of Biological Chemistry, (22 Apr 2005) Vol. 280, No. 16, pp. 16045-16052.

Refs: 51

ISSN: 0021-9258 CODEN: JBCHA3

- AU Jin Z.; Xin M.; Deng X.
- AN 2005206818 EMBASE
- L62 ANSWER 5 OF 280 MEDLINE on STN DUPLICATE 3
- TI Nongenotropic, anti-apoptotic signaling of lalpha,25(OH)2-vitamin D3 and analogs through the ligand binding domain of the vitamin D receptor in osteoblasts and osteocytes. Mediation by Src, phosphatidylinositol 3-, and .TNK kinases
- SO Journal of biological chemistry, (2005 Apr 8) 280 (14) 14130-7. Electronic Publication: 2005-01-25. Journal code: 2985121R. ISSN: 0021-9258.
- AU Vertino Anthony M; Bula Craig M; Chen Jin-Ran; Almeida Maria; Han Li; Bellido Teresita; Kousteni Stavroula; Norman Anthony W; Manolagas Stavros
- AN 2005173203 MEDLINE
- L62 ANSWER 6 OF 280 MEDLINE on STN DUPLICATE 4
- TI Flow shear stress stimulates Gabl tyrosine phosphorylation to mediate protein kinase B and endothelial nitric-oxide synthase activation in endothelial cells.
- SO Journal of biological chemistry, (2005 Apr 1) 280 (13) 12305-9. Electronic Publication: 2005-01-21. Journal code: 2985121R. ISSN: 0021-9258.
- AU Jin Zheng-Gen; Wong Chelsea; Wu Jie; Berk Bradford C
- AN 2005160658 MEDLINE
- L62 ANSWER 7 OF 280 MEDLINE on STN DUPLICATE 5
- TI Src kinase activity is required for integrin alphaVbeta3-mediated activation of nuclear factor-kappaB.
- SO Journal of biological chemistry, (2005 Apr 1) 280 (13) 12145-51. Electronic Publication: 2005-01-28. Journal code: 2985121R. ISSN: 0021-9258.
- AU Courter Donald L; Lomas Lucy; Scatena Marta; Giachelli Cecilia M
- AN 2005160711 MEDLINE
- L62 ANSWER 8 OF 280 MEDLINE on STN
- TI Involvement of c-Src and protein kinase C delta in the inhibition of Cl(-)/OH- exchange activity in Caco-2 cells by serotonin.
- SO Journal of biological chemistry, (2005 Mar 25) 280 (12) 11859-68. Electronic Publication: 2005-01-06. Journal code: 2985121R. ISSN: 0021-9258.
- AU Saksena Seema; Gill Ravinder K; Tyagi Sangeeta; Alrefai Waddah A; Sarwar Zaheer; Ramaswamy Krishnamurthy; Dudeja Pradeep K
- AN 2005146181 MEDLINE
- L62 ANSWER 9 OF 280 EMBASE COPYRIGHT 2005 ELSEVIER INC. ALL RIGHTS RESERVED. on STN
- TI Inhibition of Src-like kinases reveals Akt-dependent and -independent pathways in insulin-like growth factor I-mediated oligodendrocyte progenitor survival.
- SO Journal of Biological Chemistry, (11 Mar 2005) Vol. 280, No. 10, pp. 8918-8928.

Refs: 67

ISSN: 0021-9258 CODEN: JBCHA3

- AU Cui Q.-L.; Zheng W.-H.; Quirion R.; Almazan G.
- AN 2005138590 EMBASE
- L62 ANSWER 10 OF 280 EMBASE COPYRIGHT 2005 ELSEVIER INC. ALL RIGHTS RESERVED. on STN
- TI Internalization and Src activity regulate the time course of ERK activation, by delta opioid receptor ligands.
- SO Journal of Biological Chemistry, (4 Mar 2005) Vol. 280, No. 9, pp. 7808-7816. Refs: 39

ISSN: 0021-9258 CODEN: JBCHA3

- AU Audet N.; Paquin-Gobeil M.; Landry-Paquet O.; Schiller P.W.; Pineyro G.
- AN 2005118846 EMBASE
- L62 ANSWER 11 OF 280 MEDLINE on STN DUPLICATE 6
- TI Phorbol 12-myristate 13-acetate induces epidermal growth factor receptor transactivation via protein kinase Cdelta/c-Src pathways in glioblastoma cells.
- SO Journal of biological chemistry, (2005 Mar 4) 280 (9) 7729-38. Electronic Publication: 2004-12-23.

 Journal code: 2985121R. ISSN: 0021-9258.
- AU Amos Samson; Martin Patrick M; Polar Gregory A; Parsons Sarah J; Hussaini Isa M
- AN 2005104092 MEDLINE
- L62 ANSWER 12 OF 280 MEDLINE ON STN DUPLICATE 7
- TI Tyrosine 394 is phosphorylated in Alzheimer's paired helical filament tau and in fetal tau with c-Abl as the candidate tyrosine kinase.
- Journal of neuroscience: official journal of the Society for Neuroscience, (2005 Jul 13) 25 (28) 6584-93.

 Journal code: 8102140. ISSN: 1529-2401.
- AU Derkinderen Pascal; Scales Timothy M E; Hanger Diane P; Leung Kit-Yi; Byers Helen L; Ward Malcolm A; Lenz Christof; Price Caroline; Bird Ian N; Perera Timothy; Kellie Stuart; Williamson Ritchie; Noble Wendy; Van Etten Richard A; Leroy Karelle; Brion Jean-Pierre; Reynolds C Hugh; Anderton Brian H
- AN 2005360848 IN-PROCESS
- L62 ANSWER 13 OF 280 EMBASE COPYRIGHT 2005 ELSEVIER INC. ALL RIGHTS RESERVED. on STN
- TI Progestins induce transcriptional activation of signal transducer and activator of transcription 3 (Stat3) via a Jak- and Src-dependent mechanism in breast cancer cells.
- SO Molecular and Cellular Biology, (2005) Vol. 25, No. 12, pp. 4826-4840. Refs: 59
 ISSN: 0270-7306 CODEN: MCEBD4
- AU Proietti C.; Salatino M.; Rosemblit C.; Carnevale R.; Pecci A.; Kornblihtt A.R.; Molinolo A.A.; Frahm I.; Charreau E.H.; Schillaci R.; Elizalde P.V.
- AN 2005253451 EMBASE
- L62 ANSWER 14 OF 280 EMBASE COPYRIGHT 2005 ELSEVIER INC. ALL RIGHTS RESERVED. on STN
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recombinant protein production for use in drug screening and disease MERCKEN L; ZAMBRANO N; RUSSO T ΑU AN 2004-14884 BIOTECHDS EP 1413887 28 Apr 2004 ΡI ANSWER 70 OF 280 HCAPLUS COPYRIGHT 2005 ACS on STN L62 4-Amino-substituted derivatives of pyrazolo[3,4-d] TIpyrimidine and pyrrolo[2,3-d]pyrimidine and their preparation, pharmaceutical compositions, and use as antitumor and antileukemic agents SO PCT Int. Appl., 29 pp. CODEN: PIXXD2 Bondavalli, Francesco; Botta, Maurizio; Bruno, Olga; Manetti, Fabrizio; IN Schenone, Silvia; Carraro, Fabio 2004:1059356 HCAPLUS ΑN 142:38272 DN APPLICATION NO. DATE PATENT NO. KIND DATE **-** - - -WO 2004106340 A2 20041209 WO 2004-IT303 20040526 ΡĪ **A3** 20050217 WO 2004106340 AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG ANSWER 71 OF 280 HCAPLUS COPYRIGHT 2005 ACS on STN L62 Method using a Src family tyrosine kinase inhibitor ΤI for the treatment of myocardial infarction U.S. Pat. Appl. Publ., 44 pp., Cont.-in-part of Appl. No. PCT/US03/37653. SO CODEN: USXXCO Cheresh, David A.; Paul, Robert; Eliceiri, Brian ΙN 2004:905617 HCAPLUS ANDN 141:374724 APPLICATION NO. DATE DATE PATENT NO. KIND ______ _ _ _ _ PΙ US 2004214836 A1 20041028 US 2004-801050 20040315 WO 1999-US11780 WO 9961590 Α1 19991202 19990528 AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, W: DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG US 6685938 B1 20040203 US 1999-470881 19991222 20030710 US 2002-298377 20021118 US 2003130209 **A1** WO 2003-US37653 20031118 WO 2004045563 A2 20040603 A3 20041223 WO 2004045563 AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ,

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 - W: AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ DE DK DM DZ EC EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NO NZ OM PH PL PT RO RU SC SD SE SG SK SL TJ TM TN TR TT TZ UA UG US UZ VC VN YU ZA ZM ZW

AU 2003279795 A1 20040504 (200465)

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- AB New 4-aminopyrazolo[3,4-d] pyrimidines bearing various substituents at the position 1 and 6, were synthesized. The new compounds showed antiproliferative activity toward A431 cells, were found to be inhibitors of Src phosphorylation, and induced apoptotic cell death. In particular, 2h was a better inhibitor of Src phosphorylation than the reference compound PP2.

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- The cellular signaling machinery is a complex network of cross-talking AB proteins that enables dynamic communication between upstream causal factors and downstream effectors. Non-receptor tyrosine kinases, including Src, are the intermediates of signal transfer, controlling pathways as diverse as cell growth, death, differentiation, migration, and genome maintenance. When expressed as viral genes these proteins are potent carcinogens. Furthermore, analogous genetic alterations are observed, albeit not frequently, in human tumors. In a variety of tumors including those derived from the colon and breast, Src is either over expressed or constitutively active in a large percentage of patients. Increased expression or activity of Src correlates with the stage and metastatic potential of some neoplasia. The detailed knowledge of Src activation facilitates rational design of drugs that potentially interfere with either binding of ATP or substrate peptides. Several existing inhibitors are available as lead compounds for further development of Src inhibitors. . COPYRGT. 2004 Bentham Science Publishers Ltd.

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